

REMARKS

The above amendments are made to place the claims in a more traditional format.

Respectfully submitted,

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MARKED UP CLAIMS

- 5. (Amended) A variant of a polypeptide as defined in [any one of claims 1 to 4] claim 1, said variant differing from the polypeptide by the presence of from 1 to 5 amino acid substitutions in the sequence of said polypeptide, said variant being capable of antagonising the heterodimerization of a DP protein with an E2F protein.
- 7. (Amended) A polypeptide which comprises a first portion having the amino acid sequence of a polypeptide defined in [any one of claims 1 to 6] <u>claim 1</u> and a second portion, attached to the N- or C-terminus of the first portion, which comprises a sequence of amino acid not naturally contiguous to the first portion in DP-1.
- 10. (Amended) A pharmaceutical composition comprising a polypeptide according to [any one of the preceding claims] claim 1 together with a pharmaceutically acceptable diluent or carrier.
- 12. (Amended) A polypeptide according to [any one of claims 1 to 9 or a composition according to claim 10] <u>claim 1</u> for use in a method of treatment of the human or animal body.
- 13. (Amended) A method of inducing apoptosis in a cell which comprises introducing into said cell an effective amount of a polypeptide according to [any one of claims 1 to 9] claim 1.
- 16. (Amended) A product comprising a polypeptide as defined in [any one of claims 1 to 9] <u>claim 1</u> and a cytostatic or cytotoxic agent as a combined preparation for separate or sequential use in a method of treatment of the human or animal body.
 - 17. (Amended) A surgical stent which comprises a polypeptide as defined in

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[any one of claims 1 to 9] claim 1 in a pharmaceutically acceptable carrier.

18. (Amended) An expression vector comprising a promoter operably linked to a sequence encoding a polypeptide as defined in [any one of claims 1 to 9] claim 1.